

**AMENDMENTS TO THE CLAIMS**

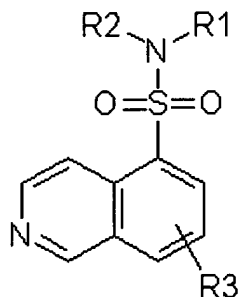
Please cancel, without prejudice, claim 18.

Please add new claims 22-26

1. **(Currently amended)** A method for ~~modulating inhibiting or reducing the growth state proliferation or growth of lung cancer cells tissue or cells derived therefrom~~, comprising contacting the ~~cells tissue~~ with an amount of an agent effective to ~~alter~~ inhibit or reduce the proliferation, ~~differentiation, or growth survival~~ of the lung cancer cells tissue, wherein the agent is ~~selected from a hedgehog therapeutic or a ptc therapeutic~~ which inhibits or attenuates hedgehog/patched signal transduction thereby inhibiting or reducing cell proliferation or growth, and wherein the agent is a small organic molecule.
2. **(Currently amended)** A method for inducing the formation of, or the maintenance or functional performance of normal lung tissue, comprising contacting the lung tissue with an amount of an agent effective to induce the formation of new lung tissue, wherein the agent is ~~selected from a hedgehog therapeutic or a ptc therapeutic~~ that promotes hedgehog/patched signal transduction thereby inducing the formation of, or the maintenance or functional performance of normal lung tissue.
3. **(Currently amended)** The method of claim 1, wherein the lung ~~tissue~~ cancer cell is in culture, and the agent is provided as a cell culture additive.
4. **(Original)** The method of claim 1, wherein the cell is treated in an animal and the agent is administered to the animal as a therapeutic composition.
5. **(Withdrawn)** The method of claim 1, wherein the agent is a hedgehog therapeutic.
6. **(Withdrawn)** The method of claim 5, wherein the hedgehog therapeutic is a polypeptide including a hedgehog polypeptide sequence of at least a bioactive extracellular portion of a hedgehog protein.

7. **(Withdrawn)** The method of claim 6, wherein the polypeptide includes at least 50 amino acids residues of an N-terminal half of the hedgehog protein.
8. **(Withdrawn)** The method of claim 6, wherein the polypeptide includes at least 100 amino acids of an extracellular domain of the hedgehog protein.
9. **(Withdrawn)** The method of claim 6, wherein the polypeptide includes at least a portion of the hedgehog protein corresponding to a 19kd fragment of an extracellular domain of the hedgehog protein.
10. **(Withdrawn)** The method of claim 6, wherein the hedgehog protein is encoded by a gene of a vertebrate organism.
11. **(Withdrawn)** The method of claim 6, wherein the polypeptide includes a hedgehog polypeptide sequence represented in the general formula of SEQ ID No. 21.
12. **(Withdrawn)** The method of claim 6, wherein the polypeptide includes a hedgehog polypeptide sequence represented in the general formula of SEQ ID No. 22.
13. **(Withdrawn)** The method of claim 6, wherein the hedgehog protein is encoded by a human hedgehog gene.
14. **(Withdrawn)** The method of claim 6, wherein the hedgehog polypeptide sequence is at least 60 percent identical to an amino acid sequence of a hedgehog protein selected from SEQ ID No:9, SEQ ID No:10, SEQ ID No:11, SEQ ID No:12, SEQ ID No:13, SEQ ID No:14, SEQ ID No:15, or SEQ ID No:16.
15. **(Withdrawn)** The method of claim 6, wherein the hedgehog polypeptide sequence is encodable by a nucleotide sequence which hybridizes under stringent conditions to a sequence selected from SEQ ID No:1, SEQ ID No:2, SEQ ID No:3, SEQ ID No:4, SEQ ID No:5, SEQ ID No:6, SEQ ID No:7, or SEQ ID No:8.

16. **(Withdrawn)** The method of claim 6, wherein the hedgehog polypeptide sequence is an amino acid sequence of a hedgehog protein selected from SEQ ID No:9, SEQ ID No:10, SEQ ID No:11, SEQ ID No:12, SEQ ID No:13, SEQ ID No:14, SEQ ID No:15, or SEQ ID No:16.
17. **(Withdrawn)** The method of claim 6, wherein the hedgehog polypeptide sequence is an amino acid sequence of a Sonic hedgehog protein.
18. **(Cancelled)**
19. **(Currently amended)** The method of claim ~~18~~2, wherein the ptc therapeutic is a small organic molecule which binds to a patched protein and derepresses patched-mediated inhibition of mitosis.
20. **(Currently amended)** The method of claim ~~18~~2, wherein the ptc therapeutic binds to patched and mimics hedgehog, thereby promoting hedgehog/patched ~~-mediated patched~~-signal transduction.
21. **(Original)** The method of claim 20, wherein the ptc therapeutic is a small organic molecule.
22. **(New)** The method of claim 2, wherein the ptc therapeutic is a protein kinase A inhibitor or a ptc antisense construct.
23. **(New)** The method of claim 22, wherein the protein kinase A inhibitor is selected from N-[2-((p-bromocinnamyl)amino)ethyl]-5-isoquinolinesulfonamide, 1-(5-isoquinoline-sulfonyl)-2-methylpiperazine, KT5720, 8-bromo-cAMP, dibutyryl-cAMP and PKA Heat Stable Inhibitor isoform  $\alpha$ .
24. **(New)** The method of claim 22, wherein the protein kinase A inhibitor is selected from a molecule of Formula I



wherein  $\text{R}_1$  and  $\text{R}_2$  each can independently represent hydrogen, and as valence and stability permit a lower alkyl, a lower alkenyl, a lower alkynyl, a carbonyl (such as a carboxyl, an ester, a formate, or a ketone), a thiocarbonyl (such as a thioester, a thioacetate, or a thioformate), an amino, an acylamino, an amido, a cyano, a nitro, an azido, a sulfate, a sulfonate, a sulfonamido,  $-(\text{CH}_2)_m\text{R}_8$ ,  $-(\text{CH}_2)_m\text{OH}$ ,  $-(\text{CH}_2)_m\text{O-lower alkyl}$ ,  $-(\text{CH}_2)_m\text{O-lower alkenyl}$ ,  $-(\text{CH}_2)_n\text{O}-(\text{CH}_2)_m\text{R}_8$ ,  $-(\text{CH}_2)_m\text{SH}$ ,  $-(\text{CH}_2)_m\text{S-lower alkyl}$ ,  $-(\text{CH}_2)_m\text{S-lower alkenyl}$ ,  $-(\text{CH}_2)_n\text{S}-(\text{CH}_2)_m\text{R}_8$ , or

$\text{R}_1$  and  $\text{R}_2$  taken together with N form a heterocycle (substituted or unsubstituted);

$\text{R}_3$  is absent or represents one or more substitutions to the isoquinoline ring such as a lower alkyl, a lower alkenyl, a lower alkynyl, a carbonyl (such as a carboxyl, an ester, a formate, or a ketone), a thiocarbonyl (such as a thioester, a thioacetate, or a thioformate), an amino, an acylamino, an amido, a cyano, a nitro, an azido, a sulfate, a sulfonate, a sulfonamido,  $-(\text{CH}_2)_m\text{R}_8$ ,  $-(\text{CH}_2)_m\text{OH}$ ,  $-(\text{CH}_2)_m\text{O-lower alkyl}$ ,  $-(\text{CH}_2)_m\text{O-lower alkenyl}$ ,  $-(\text{CH}_2)_n\text{O}-(\text{CH}_2)_m\text{R}_8$ ,  $-(\text{CH}_2)_m\text{SH}$ ,  $-(\text{CH}_2)_m\text{S-lower alkyl}$ ,  $-(\text{CH}_2)_m\text{S-lower alkenyl}$ ,  $-(\text{CH}_2)_n\text{S}-(\text{CH}_2)_m\text{R}_8$ ;

$\text{R}_8$  represents a substituted or unsubstituted aryl, aralkyl, cycloalkyl, cycloalkenyl, or heterocycle; and

n and m are independently for each occurrence zero or an integer in the range of 1 to 6.

25. (New) The method of claim 1, wherein the lung cancer cells are small cell lung cancer (SCLC) cells or non-small cell lung cancer (NSCLC) cells.

26. (New) The method of claim 1, wherein the lung cancer cells are adenocarcinoma cells, lung cell carcinoma cells, or squamous cell carcinoma cells.